FILE 'HOME' ENTERED AT 16:36:46 ON 19 OCT 2000

=> fil .bec

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

0 15

0.15

FULL ESTIMATED COST

FILES 'MEDLINE, SCISEARCH, LIFESCI, BIOTECHDS, BIOSIS, EMBASE, HCAPLUS, NTIS, ESBIOBASE, BIOTECHNO' ENTERED AT 16:36:55 ON 19 OCT 2000 ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.

10 FILES IN THE FILE LIST

=> s sulfolobus or acidocaldarius

FILE 'MEDLINE'

770 SULFOLOBUS

342 ACIDOCALDARIUS .

L1 838 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'SCISEARCH'

1326 SULFOLOBUS

730 ACIDOCALDARIUS

L2 1463 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'LIFESCI'

705 SULFOLOBUS

344 ACIDOCALDARIUS

L3 772 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'BIOTECHDS'

312 SULFOLOBUS

163 ACIDOCALDARIUS

L4 363 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'BIOSIS'

1201 SULFOLOBUS

647 ACIDOCALDARIUS

L5 1357 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'EMBASE'

728 SULFOLOBUS

343 ACIDOCALDARIUS

L6 806 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'HCAPLUS'

1353 SULFOLOBUS

678 ACIDOCALDARIUS

L7 1507 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'NTIS'

37 SULFOLOBUS

14 ACIDOCALDARIUS

L8 40 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'ESBIOBASE'

454 SULFOLOBUS

181 ACIDOCALDARIUS

L9 491 SULFOLOBUS OR ACIDOCALDARIUS

FILE 'BIOTECHNO'

658 SULFOLOBUS

275 ACIDOCALDARIUS

709 SULFOLOBUS OR ACIDOCALDARIUS L10

TOTAL FOR ALL FILES

8346 SULFOLOBUS OR ACIDOCALDARIUS

=> s trehalose

FILE 'MEDLINE'

L12 1972 TREHALOSE

FILE 'SCISEARCH'

L13 2535 TREHALOSE

FILE 'LIFESCI'

L14 1308 TREHALOSE

FILE 'BIOTECHDS'

L15 . 521 TREHALOSE

FILE 'BIOSIS'

L16 3823 TREHALOSE

FILE 'EMBASE'

L17 1935 TREHALOSE

FILE 'HCAPLUS'

L18 5656 TREHALOSE

FILE 'NTIS'

L19 61 TREHALOSE

FILE 'ESBIOBASE'

L20 805 TREHALOSE

FILE 'BIOTECHNO'

L21 1037 TREHALOSE

TOTAL FOR ALL FILES

19653 TREHALOSE

=> s non-reducing saccharide#

FILE 'MEDLINE'

2433763 NON

73955 REDUCING

2623 SACCHARIDE#

1 NON-REDUCING SACCHARIDE#

(NON (W) REDUCING (W) SACCHARIDE#)

FILE 'SCISEARCH'

518884 NON

82111 REDUCING

3553 SACCHARIDE#

```
2 NON-REDUCING SACCHARIDE#
L24
                (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'LIFESCI'
       118756 "NON"
        19077 "REDUCING"
         1005 SACCHARIDE#
          3 NON-REDUCING SACCHARIDE#
L25
                ("NON"(W) "REDUCING"(W) SACCHARIDE#)
FILE 'BIOTECHDS'
       22327 NON
         5859 REDUCING
          706 SACCHARIDE#
           15 NON-REDUCING SACCHARIDE#
               (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'BIOSIS'
  474771 NON
        80070 REDUCING
         27137 SACCHARIDE#
L27
           13 NON-REDUCING SACCHARIDE#
                (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'EMBASE'
        428771 "NON"
        70859 "REDUCING"
         2332 SACCHARIDE#
            0 NON-REDUCING SACCHARIDE#
               ("NON"(W) "REDUCING"(W) SACCHARIDE#)
FILE 'HCAPLUS'
        410977 NON
        186467 REDUCING
       10184 SACCHARIDE#
           19 NON-REDUCING SACCHARIDE#
                (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'NTIS'
         84456 NON
         24538 REDUCING
           250 SACCHARIDE#
             0 NON-REDUCING SACCHARIDE#
L30
               (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'ESBIOBASE'
       127286 NON
        19949 REDUCING
           810 SACCHARIDE#
            0 NON-REDUCING SACCHARIDE#
L31
                (NON (W) REDUCING (W) SACCHARIDE#)
FILE 'BIOTECHNO'
         95655 NON
       15581 REDUCING
           934 SACCHARIDE#
             0 NON-REDUCING SACCHARIDE#
L32
                 (NON(W) REDUCING(W) SACCHARIDE#)
```

TOTAL FOR ALL FILES

```
53 NON-REDUCING SACCHARIDE#
L33
=> s (122 or 133)(8a)(synthes? or produc? or form######)(5a)enzym?
FILE 'MEDLINE'
        360157 SYNTHES?
        892273 PRODUC?
        925210 FORM######
        854146 ENZYM?
            39 (L12 OR L23) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L34
FILE 'SCISEARCH'
       607484 SYNTHES?
       1148375 PRODUC?
       1300302 FORM######
        369456 ENZYM?
            51 (L13 OR L24)(8A)(SYNTHES? OR PRODUC? OR FORM######)(5A)ENZYM?
L35
FILE 'LIFESCI'
        111542 SYNTHES?
        378068 PRODUC?
        287187 FORM######
       174869 ENZYM?
            36 (L14 OR L25) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L36
FILE 'BIOTECHDS'
       20899 SYNTHES?
        148825 PRODUC?
         59860 FORM######
         87824 ENZYM?
            48 (L15 OR L26) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L37
FILE 'BIOSIS'
       519392 SYNTHES?
       1257082 PRODUC?
       1094999 FORM######
        909774 ENZYM?
            70 (L16 OR L27) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L38
FILE 'EMBASE'
        449294 SYNTHES?
        895958 PRODUC?
        830949 FORM######
        591498 ENZYM?
            31 (L17 OR L28)(8A)(SYNTHES? OR PRODUC? OR FORM######)(5A)ENZYM?
L39
FILE 'HCAPLUS'
       .999290 SYNTHES?
       2470442 PRODUC?
       625169 PRODN
       2781937 PRODUC?
                 (PRODUC? OR PRODN)
       3685888 FORM######
        759973 ENZYM?
           119 (L18 OR L29) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L40
FILE 'NTIS'
        39512 SYNTHES?
        341830 PRODUC?
```

269806 FORM######

```
11891 ENZYM?
         4 (L19 OR L30) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
FILE 'ESBIOBASE'
       105954 SYNTHES?
       287167 PRODUC?
       233496 FORM######
       146854 ENZYM?
           26 (L20 OR L31) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
FILE 'BIOTECHNO'
       135351 SYNTHES?
       298474 PRODUC?
       238674 FORM######
       270706 ENZYM?
           32 (L21 OR L32) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
L43
TOTAL FOR ALL FILES
         456 (L22 OR L33) (8A) (SYNTHES? OR PRODUC? OR FORM######) (5A) ENZYM?
=> s 144 and thermostab?
FILE 'MEDLINE'
     4753 THERMOSTAB?
          3 L34 AND THERMOSTAB?
FILE 'SCISEARCH'
        6070 THERMOSTAB?
         7 L35 AND THERMOSTAB?
FILE 'LIFESCI'
         2930 THERMOSTAB?
            5 L36 AND THERMOSTAB?
FILE 'BIOTECHDS'
         5495 THERMOSTAB?
          13 L37 AND THERMOSTAB?
FILE 'BIOSIS'
         8021 THERMOSTAB?
         9 L38 AND THERMOSTAB?
FILE 'EMBASE'
     7587 THERMOSTAB?
          5 L39 AND THERMOSTAB?
FILE 'HCAPLUS'
        13309 THERMOSTAB?
          11 L40 AND THERMOSTAB?
FILE 'NTIS'
          178 THERMOSTAB?
           0 L41 AND THERMOSTAB?
L52
FILE 'ESBIOBASE'
         1943 THERMOSTAB?
           3 L42 AND THERMOSTAB?
L53
```

FILE 'BIOTECHNO'

4641 THERMOSTAB?

TOTAL FOR ALL FILES

L55 61 L44 AND THERMOSTAB?

=> s 111 and (122 or 133)

FILE 'MEDLINE'

L56 11 L1 AND (L12 OR L23)

FILE 'SCISEARCH'

L57 23 L2 AND (L13 OR L24)

FILE 'LIFESCI'

L58 9 L3 AND (L14 OR L25)

FILE 'BIOTECHDS'

L59 19 L4 AND (L15 OR L26)

FILE 'BIOSIS'

L60 17 L5 AND (L16 OR L27)

FILE 'EMBASE'

L61 5 L6 AND (L17 OR L28)

FILE 'HCAPLUS'

L62 32 L7 AND (L18 OR L29)

FILE 'NTIS'

L63 0 L8 AND (L19 OR L30)

FILE 'ESBIOBASE'

L64 12 L9 AND (L20 OR L31)

FILE 'BIOTECHNO'

L65 12 L10 AND (L21 OR L32)

TOTAL FOR ALL FILES

L66 140 L11 AND (L22 OR L33)

=> s (155 or 166)not 1996/py range=,1996

FILE 'MEDLINE'

223309 1996/PY

L67 0 (L45 OR L56)NOT 1996/PY

FILE 'SCISEARCH'

794148 1996/PY

L68 0 (L46 OR L57)NOT 1996/PY

FILE 'LIFESCI'

43658 1996/PY

L69 1 (L47 OR L58) NOT 1996/PY

FILE 'BIOTECHDS'

11161 1996/PY

L70 6 (L48 OR L59) NOT 1996/PY

FILE 'BIOSIS'

405433 1996/PY

```
1 (L49 OR L60)NOT 1996/PY
 L71
 FILE 'EMBASE'
        338946 1996/PY
             0 (L50 OR L61)NOT 1996/PY
 FILE 'HCAPLUS'
       659369 1996/PY
            3 (L51 OR L62)NOT 1996/PY
L73
 FILE 'NTIS'
       14160 1996/PY
            0 (L52 OR L63)NOT 1996/PY
 FILE 'ESBIOBASE'
        151367 1996/PY
            0 (L53 OR L64)NOT 1996/PY
 FILE 'BIOTECHNO'
       0 1996/PY
             0 (L54 OR L65)NOT 1996/PY
 TOTAL FOR ALL FILES
 L77 11 (L55 OR L66) NOT 1996/PY
 => s (155 or 166) and py=<1995 range=1997,
 FILE 'MEDLINE'
        18926 PY=<1995
       0 (L45 OR L56) AND PY=<1995
 FILE 'SCISEARCH'
         525 PY=<1995
           0 (L46 OR L57) AND PY=<1995
 FILE 'LIFESCI'
       13245 PY=<1995
          0 (L47 OR L58) AND PY=<1995
 FILE 'BIOTECHDS'
           255 PY=<1995
                (PY = < 1995)
 L81
            0 (L48 OR L59) AND PY=<1995
 FILE 'BIOSIS'
          5732 PY=<1995
 L82
          0 (L49 OR L60) AND PY=<1995
 FILE 'EMBASE'
           634 PY=<1995
 L83
           0 (L50 OR L61) AND PY=<1995
 FILE 'HCAPLUS'
         33103 PY=<1995
           0 (L51 OR L62) AND PY=<1995
 FILE 'NTIS'
         64177 PY=<1995
 L85
            0 (L52 OR L63) AND PY=<1995
```

```
FILE 'ESBIOBASE'
           384 PY=<1995
             0 (L53 OR L64) AND PY=<1995
L86
FILE 'BIOTECHNO'
        845083 PY=<1995
             0 (L54 OR L65) AND PY=<1995
L87
TOTAL FOR ALL FILES
             0 (L55 OR L66) AND PY=<1995
=> dup rem 177
PROCESSING COMPLETED FOR L77
              7 DUP REM L77 (4 DUPLICATES REMOVED)
L89
=> d tot
     ANSWER 1 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD
L89
      Novel transferase and amylase production and use;
TΤ
         enzyme preparation for oligosaccharide and alpha, alpha-
       trehalose production
      Kato M; Miura Y; Kettoku M; Kobayashi K; Iwamatsu A; Komeda T
AU
AN
      1996-02920 BIOTECHDS
      WO 9534642 21 Dec 1995
PΙ
      ANSWER 2 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD
      Thermostable non-reducing
    saccharide-forming enzyme;
         non-reducing partial starch hydrolyzate or trehalose
       production using new Sulfolobus sp. enzyme
         and glucoamylase or alpha-glucosidase for use as a sweetener, etc.
ΑU
      Nakada T; Chaen H; Sugimoto T; Miyake T
AN
      1996-03026 BIOTECHDS
      EP 688867 27 Dec 1995
PΙ
      ANSWER 3 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD
L89
TΙ
      Thermostable trehalose-releasing enzyme;
         Sulfobolus acidocaldarius and Sulfobolus solfataricus
         thermostable enzyme production and characterization,
AU
      Ikegami S; Kubota M; Sugimoto T; Miyake T
AN
      1996-04132 BIOTECHDS
      EP 688866 27 Dec 1995
PΙ
      ANSWER 4 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD
L89
      Non-reducing saccharide-forming
TТ
    enzyme and its production and application;
         Arthrobacter sp. and Rhizobium sp. fermentation and enzyme
         use in alpha-glucosyl trehalose production
ΑN
      1994-11285 BIOTECHDS
      EP 606753 20 Jul 1994
PΤ
L89
      ANSWER 5 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD
TI
      Thermostable amylolytic activity from Sulfolobus
      solfataricus;
         amylase production, purification and characterization; starch
         saccharification to glucose and trehalose
SO
      Biotech Forum Eur.; (1991) 8, 4, 201-03
      Lama L; Nicolaus B; Trincone A; Morzillo P; Calandrelli V; Gambacorta A
ÀU
```

AN 1991-08311 BIOTECHDS

L89 ANSWER 6 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD Starch conversion with immobilized thermophilic archaebacterium Sulfolobus solfataricus;

glucose production from starch saccharification by thermostable glucoamylase

SO Biotechnol.Lett.; (1990) 12, 6, 431-32

CODEN: BILED3

AU Lama L; Nicolaus B; Trincone A; Morzillo P; De Rosa M; Gambacorta A

AN 1990-10306 BIOTECHDS

L89 ANSWER 7 OF 7 LIFESCI COPYRIGHT 2000 CSA DUPLICATE 3

TI Trehalose in archaebacteria.

SO SYST. APPL. MICROBIOL., (1988) vol. 10, no. 3, pp. 215-217.

AU Nicolaus, B.; Gambacorta, A.; Basso, A.L.; Riccio, R.; De Rosa, M.; Grant, W.D.

AN 88:92917 LIFESCI

=> d ab tot

ANSWER 1 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD L89 A transferase is claimed, which acts on a saccharide composed of at least AB 3 sugar units, where at least 3 glucose residues are linked alpha-1,4% so as to transfer the alpha-1,4 linkages to alpha-1,alpha-1 linkages. Also claimed are: (a) a gene coding for the transferase; (b) a process for producing an oligosaccharide using the transferase; (c) an amylase that has principal activity of acting on a saccharide composed of 3 sugar units, where at least 3 sugar units on the reducing end side are glucose units and the linkage between the first and second glucose units is alpha-1, alpha-1, while the linkage between the second and the third glucose units is alpha-1,4, so as to liberate alpha, alphatrehalose by hydrolyzing the alpha-1,4 linkage within the molecular chain of the substrate and that liberates disaccharides and monosaccharides as the principle final products; (d) a process for producing the amylase; (e) a gene coding for the amylase; and (e) a process of producing alpha-alpha-trehalose from e.g. malto-oligosaccharides by using a combination of the transferase and the amylase. (306pp)

L89 ANSWER 2 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD AB A new thermostable enzyme forms a

non-reducing saccharide with a

trehalose structure when reacted with a reducing partial starch hydrolyzate with a degree of glucose polymerization of at least 3, preferably with a trehalose end unit. The enzyme is stable up to 85 deg at pH 7.0 for 60 min, and is from a Sulfolobus sp. The enzyme has a mol.wt. of 69,000-79,000 (SDS-PAGE), a pI of 5.4-6.4, an optimum temp. of 75 deg (at pH 5.5 for 60 min), an optimum pH of 5.0-5.5(at 60 deg for 60 min), and pH stability at pH 4.0-9.5 (at 25 deg for 16 The product may be purified further by strongly acidic cation-exchange chromatography. A food product, cosmetic or pharmaceutical containing the product is also new. Trehalose (in hydrous or anhydrous crystalline form) may be produced by treating the product with glucoamylase (EC-3.2.1.3) or alpha-glucosidase (EC-3.2.1.20). The product may be used as a sweetener, osmoregulator, excipient, etc., and has higher stability than reducing starch hydrolyzates. The process may be carried out on an industrial scale at

ANSWER 3 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD

The following are claimed: (1) a thermostable trehalose

-releasing enzyme (I) which specifically hydrolyzes the linkage between the trehalose portion and the remaining glycosyl portion of a non-reducing saccharide having a terminal trehalose unit and a degree of glucose polymerization of at least

3; (2) production of (I) by culturing an (I)-producing microorganism in nutrient medium and recovering (I) from the culture; (3) a process for producing trehalose by allowing (I) to act on a solution

3; (2) production of (I) by culturing an (I)-producing microorganism in a nutrient medium and recovering (I) from the culture; (3) a process for producing trehalose by allowing (I) to act on a solution containing a saccharide as above; (4) a trehalose prepared by the above process; and (5) a composition containing the trehalose of (I), especially a food product, cosmetic or pharmaceutical. Preferably, (I) is derived from Sulfobolus acidocaldarius ATCC 33909, S. acidocaldarius ATCC 49426, Sulfobolus solfataricus ATCC 35091 and S. solfataricus ATCC 35092. (I) has better thermal stability than similar enzymes derived from Rhizobium and sp. Arthrobacter sp., and it is stable at temp. levels above 55 deg. (I) has a mol.wt. of 54,000-64,000 (SDS-PAGE), a pI of 5.6-6.6, and an optimum temp. of 75 deg when incubated at pH 6 for 30 min. (36pp)

L89 ANSWER 4 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD AB A new non-reducing saccharide-

forming enzyme (I) is prepared by culturing a bacterium of the genera Rhizobium, Arthrobacter, Brevibacterium, Flavobacterium, Micrococcus, Curtobacterium, Mycobacterium or Terrebacter or their mutants, especially Rhizobium sp. M-11 (FERM BP-4130) and Arthrobacter sp. Q36 (FERM BP-4316). (I) is capable of catalyzing the formation of trehalose-type sugars with a trehalose structure as an end unit from partial starch hydrolyzates of degree of glucose polymerization at least 3. Glucoamylase (EC-3.2.1.3) and alpha-glucosidase (EC-3.2.1.20) may then be used to convert the product to trehalose. The trehalose-type sugars are useful in food, cosmetics and pharmaceuticals. (I) has mol.wt. 76,000-87,000 (SDS-PAGE), isoelectric point 3.6 +/- 4.6 using an ampholyte, optimal activity at 35-40 deg and pH 6.4-7.2,

thermostability up to 35-40 deg at pH 7.0 for 1 hr; and pH
 stability of 5.5-11.0 at 25 deg for 16 hr. (I) forms alpha-glucosyl
 trehalose of formula Gn-T (where G = glucose residue, n = integer and T =
 alpha,alpha-trehalose). Alpha-glucosyl trehalose and its compositions
 are also new. (42pp)

ANSWER 5 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD L89 AΒ Sulfolobus solfataricus MT-4 (ATCC 49155), an archaebacterium capable of converting starch to glucose and trehalose, was grown at 87 deg in a defined culture medium. Amylase activity was isolated from the cytoplasmic fraction of disrupted cells by ammonium sulfate precipitation, acetone precipitation and dialysis. The partially purified enzyme (PPE) was stable after 5 hr at 80 deg, and showed a half-life of 1 hr at 90 deg. Amylolytic activity was detected at pH 3.5-9.0 and 30-100 deg. Optimal activity was shown at 70 deg and pH 5.5. The PPE was inhibited by Cu2+ and Zn2+, but not by Mg2+, dithiothreitol or beta-cyclodextrin. Ca2+ was not required for activity. PPE hydrolyzed starch and amylopectin to form glucose, and hydrolyzed linear oligosaccharides (more than 3 sugar units) to form glucose and malto derivatives. Non-reducing alpha, alpha'-trehalose (a potential sweetener, cosmetic and cryoprotectant) was also formed. Further enzyme purification is being investigated in order to determine whether a single enzyme is responsible for both the hydrolysis and trehalose production activities. (9 ref)

ANSWER 6 OF 7 BIOTECHDS COPYRIGHT 2000 DERWENT INFORMATION LTD L89 Alginate-immobilized Sulfolobus solfataricus strain MT-4 (ATCC AΒ 49155) was used for the saccharification of starch to produce glucose. Immobilized cells (2.5 g wet wt.) were added to starch (15 mg) in 9 ml of 10 mM acetate buffer pH 5.5 and incubated at 70 deg. Amylolytic activity of S. solfataricus converted starch to glucose and trehalose after 1 hr and 3 hr, respectively. The relative molar ratio of glucose and trehalose was 2:1. Glucose and trehalose represented 44% of the initial amount of starch. As glucose was the first product of starch hydrolysis with whole cells of S. solfataricus, the amylolytic activity was determined as a glucoamylase (EC-3.2.1.3). The formation of the non-reducing sugar trehalose, and the absence of maltose, maltotriose, etc., in the hydrolysis process suggested that a new type of enzymic activity, or multiple enzyme pathway was present in S. solfataricus. (7 ref)

ANSWER 7 OF 7 LIFESCI COPYRIGHT 2000 CSA DUPLICATE 3

The non-reducing disaccharide trehalose (alpha
-D-glucopyranosyl-alpha-D-glucopyranoside) was identified in sulfolobus solfataricus by super(13)C NMR spectroscopy. The screening of a range of other archaebacteria, using a rapid isolation and purification procedure for trehalose, indicated that this disaccharide is present in a number of halophilic archaebacteria, thermophilic and sulphur-dependent archaebacteria and methanogenic archaebacteria.

=> .log y

COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST
34.86
35.01

STN INTERNATIONAL LOGOFF AT 16:50:02 ON 19 OCT 2000